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II. AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the

application:

Claim 1 (currently amended) A method for the prevention of the loss of visual

acuity associated with AMD, which comprises, juxtasclerally administering

pharmaceutically effective amount of the compound anecortave acetate or its corresponding

alcohol, wherein said administering is by a method selected from the group consisting of

posterior juxtascleral injection, juxtascleral implant, intravitreal injection, or implant.

Claim 2 (original) The method of claim 1, wherein the compound is administered

as a juxtascleral depot.

Claim 3 (original) The method of claim 2, wherein the depot comprises 3 mg - 30

mg of compound.

The method of claim 3, wherein the depot comprises 15 mg of Claim 4 (original)

compound.

Claim 5 (currently amended) A method for maintaining visual acuity in a person

suffering from AMD, which comprises juxtasclerally administering a pharmaceutically

effective amount of the compound anecortave acetate or its corresponding alcohol, wherein

said administering is by a method selected from the group consisting of posterior juxtascleral

injection, juxtascleral implant, intravitreal injection, or implant.

Claim 6 (original) The method of claim 5, wherein the compound is administered

as a juxtascleral depot.

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Claim 7 (original) The method of claim 6, wherein the depot comprises 3 mg - 30 mg of compound.

Claim 8 (original) The method of claim 7, wherein the depot comprises 15 mg of compound.

Claim 9 (currently amended) A method for the inhibition of lesion growth associated with AMD, which comprises juxtasclerally administering a pharmaceutically effective amount of the compound anecortave acetate or its corresponding alcohol, wherein said administering is by a method selected from the group consisting of posterior juxtascleral injection, juxtascleral implant, intravitreal injection, or implant.

Claim 10 (original) The method of claim 9, wherein the compound is administered as a juxtascleral depot.

Claim 11 (original) The method of claim 10, wherein the depot comprises 3 mg – 30 mg of compound.

Claim 12 (original) The method of claim 11, wherein the depot comprises 15 mg of compound.

Claim 13 (original) The method of claim any one of claims 1, 5, or 9, wherein the compound is administered in a juxtascleral implant.

Claim 14 (new) A method for inhibiting blood vessel growth associated with AMD, said method comprising administering a pharmaceutically effective amount of the compound anecortave acetate or its accompanying alcohol, wherein the administering is by juxtascleral injection, intravitreal injection, juxtascleral implant, or other implant.

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Claim 15 (new) The method of claim 14, wherein the amount of compound administered is from 3 mg to 30 mg.

Claim 16 (new) The method of claim 15, wherein the amount of compound administered is 15 mg.

Claim 17 (new) The method of claim 9, wherein the lesion is a predominantly classic subfoveal lesion.

Claim 18 (new) The method of claim 9, wherein the lesion is a minimally classic lesion.

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III. REMARKS

A. Status of the Claims

Claims 1-13 were originally filed with the case on June 26, 2003. No Office Action

has issued in the present case. Claims 1, 5, 9 and 13 are amended herein to clarify the subject

matter of the invention. Claims 14-18 are added herein. Support for the amendments can be

found throughout the specification and in the claims as originally filed. More specifically,

support for amendments to claims 1, 5 and 9 and for new claim 14 can be found at page 10.

lines 34-36. Support for new claims 14-16 can also be found at page 1, lines 28-31 and in the

claims as originally filed. Support for new claims 17 and 18 can be found at page 6, lines 4-

6.

В. Conclusion

Applicants respectfully request that the claims be considered as amended herein.

The Examiner is invited to contact the undersigned attorney at (817) 551-4321 with

any questions, comments or suggestions relating to the referenced patent application.

Respectfully submitted,

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Date: